abcam

Product datasheet

Alexa Fluor® 488 Anti-MCM7/PRL antibody [EP1974Y] ab199753



重组 RabMAb

2 图像

概述

产品名称 Alexa Fluor® 488荧光Anti-MCM7/PRL抗体[EP1974Y]

描述 Alexa Fluor® 488荧光兔单克隆抗体[EP1974Y] to MCM7/PRL

宿主 Rabbit

偶联物 Alexa Fluor® 488. Ex: 495nm, Em: 519nm

经测试应用 适用干: ICC/IF 种属反应性 与反应: Human

预测可用于: Mouse, Rat 📤

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

阳性对照 ICC/IF: MCF7 cells.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit 常规说明

monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

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outlicensing@thermofisher.com.

性能

形式 Liquid

存放说明 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

Avoid freeze / thaw cycle. Store In the Dark.

存储溶液 pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: PBS, 30% Glycerol (glycerin, glycerine), 1% BSA

纯**度** Affinity purified

同种型 IgG

应用

The Abpromise guarantee Abpromise™承诺保证使用ab199753于以下的经测试应用

"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
ICC/IF		1/100.

靶标

功能 Acts as component of the MCM2-7 complex (MCM complex) which is the putative replicative

helicase essential for 'once per cell cycle' DNA replication initiation and elongation in eukaryotic cells. The active ATPase sites in the MCM2-7 ring are formed through the interaction surfaces of two neighboring subunits such that a critical structure of a conserved arginine finger motif is provided in trans relative to the ATP-binding site of the Walker A box of the adjacent subunit. The six ATPase active sites, however, are likely to contribute differentially to the complex helicase

activity. Required for S-phase checkpoint activation upon UV-induced damage.

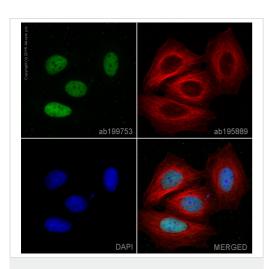
序列相似性 Belongs to the MCM family.

Contains 1 MCM domain.

翻译后修饰 Phosphorylated upon DNA damage, probably by ATM or ATR.

细胞定位 Nucleus.

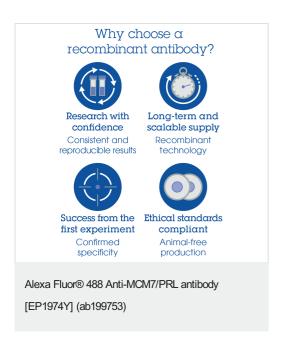
图片



Immunocytochemistry/ Immunofluorescence - Alexa Fluor® 488 Anti-MCM7/PRL antibody [EP1974Y] (ab199753)

ab199753 staining MCM7/PRL in MCF7 cells. The cells were fixed with 4% formaldehyde (10min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated overnight at +4°C with ab199753 at 1/100 dilution (shown in green) and <u>ab195889</u>, Mouse monoclonal to alpha Tubulin (Alexa Fluor® 594), at $2\mu g/ml$ (shown in red). Nuclear DNA was labelled with DAPI (shown in blue).

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).



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